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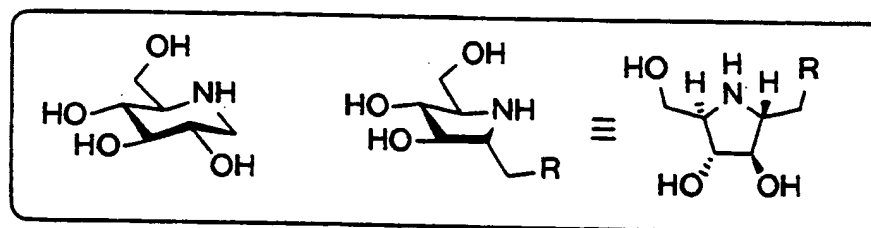
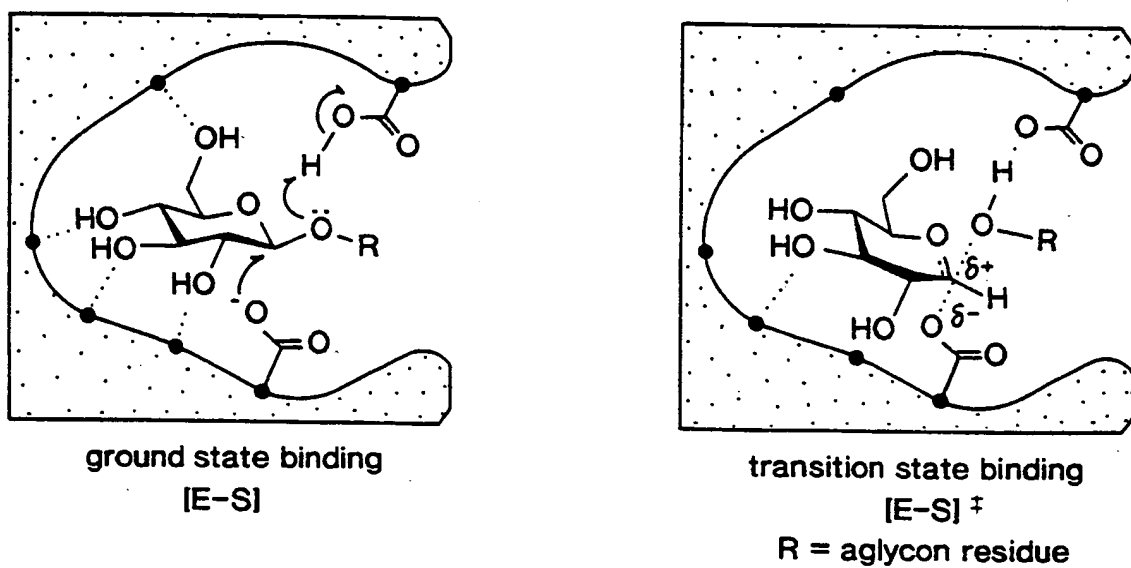


FIG. 1

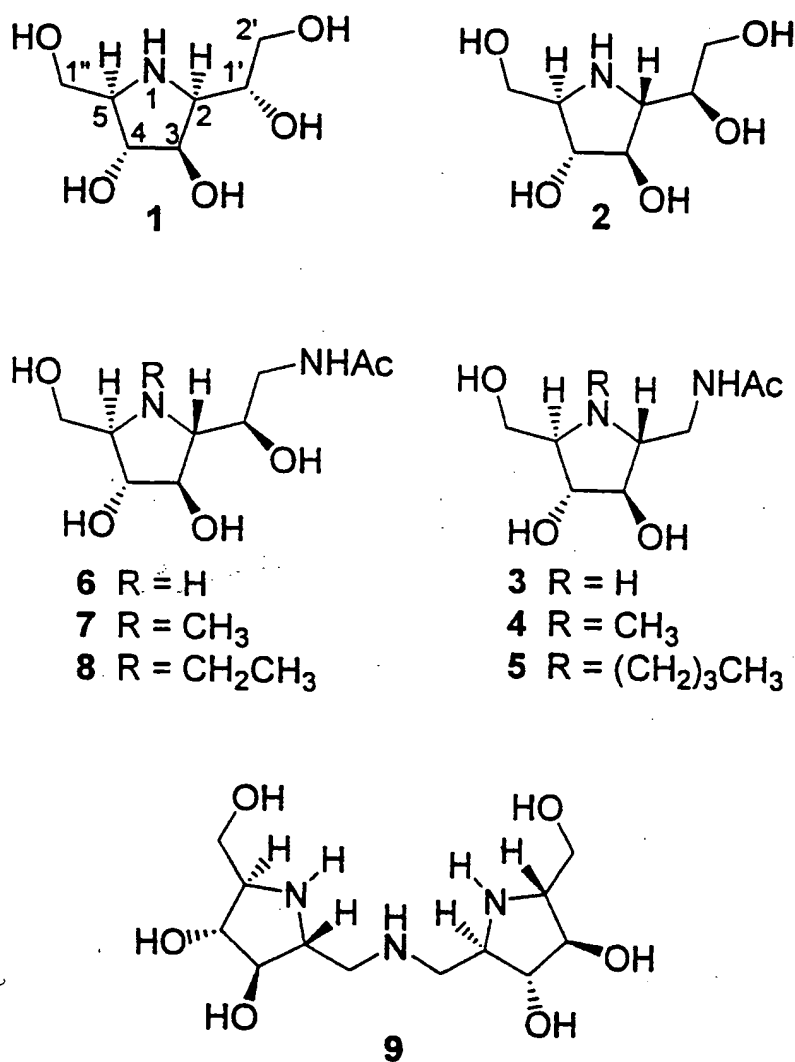


FIG. 2

K_i (μ M)					
compd	α -glucosidase ^a <i>Saccharomyces</i> sp	β -glucosidase ^b sweet almond	β -N-acetylglucosaminidase		
			bovine kidney ^c	human placenta A ^d	p ^e
1 ^f	330	50	^h	-	-
2 ^f	28	2.6	-	-	-
3	380	* ^g	2.9×10^{-1}	2.2×10^{-1}	2.6×10^{-1}
4	ni	ni	1.1×10^{-1}	1.4×10^{-1}	8.0×10^{-2}
5	ni	ni	1.3	5.1×10^{-1}	2.4×10^{-1}
6	*	2.2	*	-	-
7	*	45	*	-	-
8	ni	120	ni ⁱ	-	-
9	53	37	-	-	-

^a $K_m = 0.30$ mM, $V_{max} = 0.7$ (μ M/s)/mg, ^b $K_m = 3.2$ mM, $V_{max} = 3.2$ (μ M/s)/mg, ^c $K_m = 4.1$ mM, $V_{max} = 6.4$ (μ M/s)/mg, ^d $K_m = 2.5$ mM, $V_{max} = 2.1$ (μ M/s)/mg, ^e $K_m = 2.8$ mM, $V_{max} = 2.3$ (μ M/s)/mg, ^f Preliminary assay result using photometric assay gave K_i values: 430 and 18 μ M for compound 1 and 7.2 and 7.6 μ M for compound 2 toward α -glucosidase and β -glucosidase, respectively. See also refs 6a and 19. * *: poor inhibitor with IC_{50} above 0.5 mM. ^h -: not tested. ⁱ ni: not inhibitor.

FIG. 3

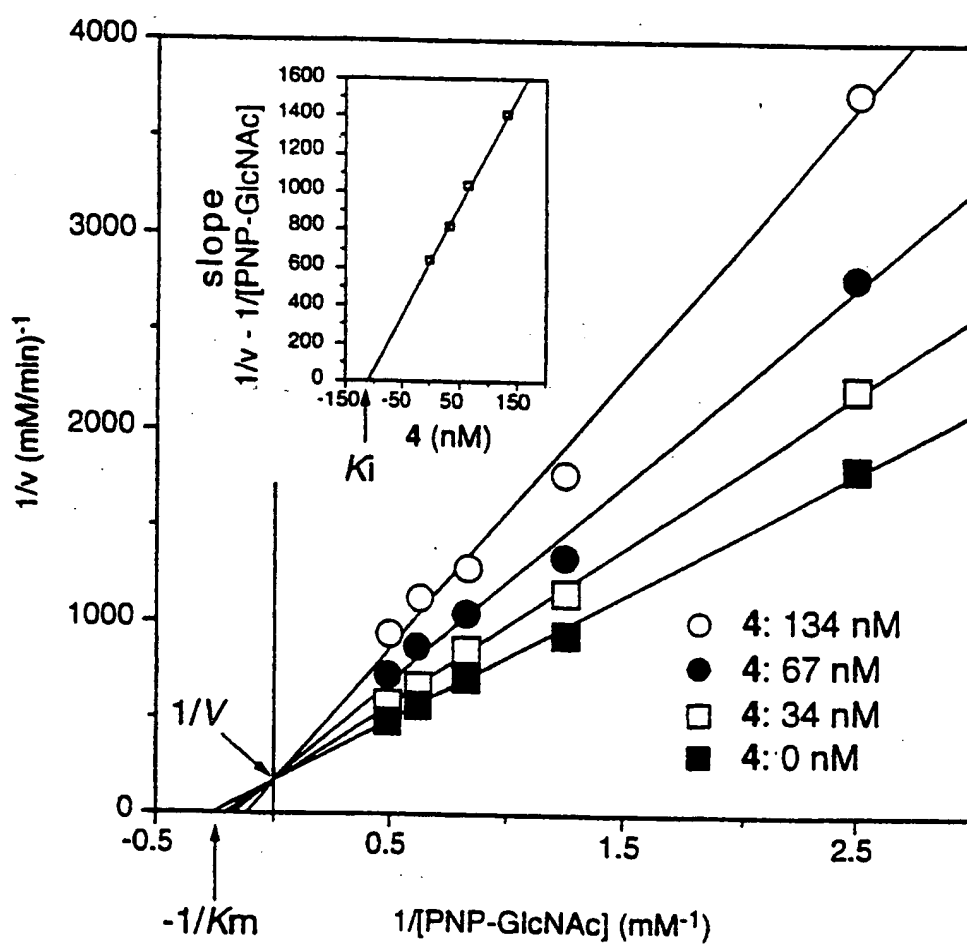
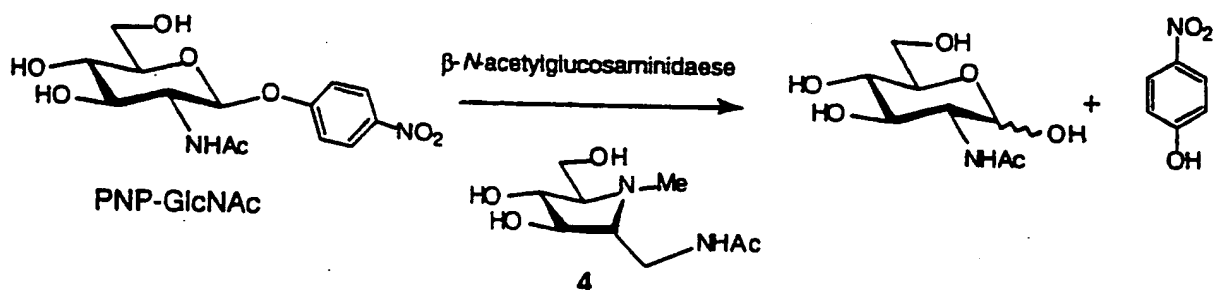


FIG. 4

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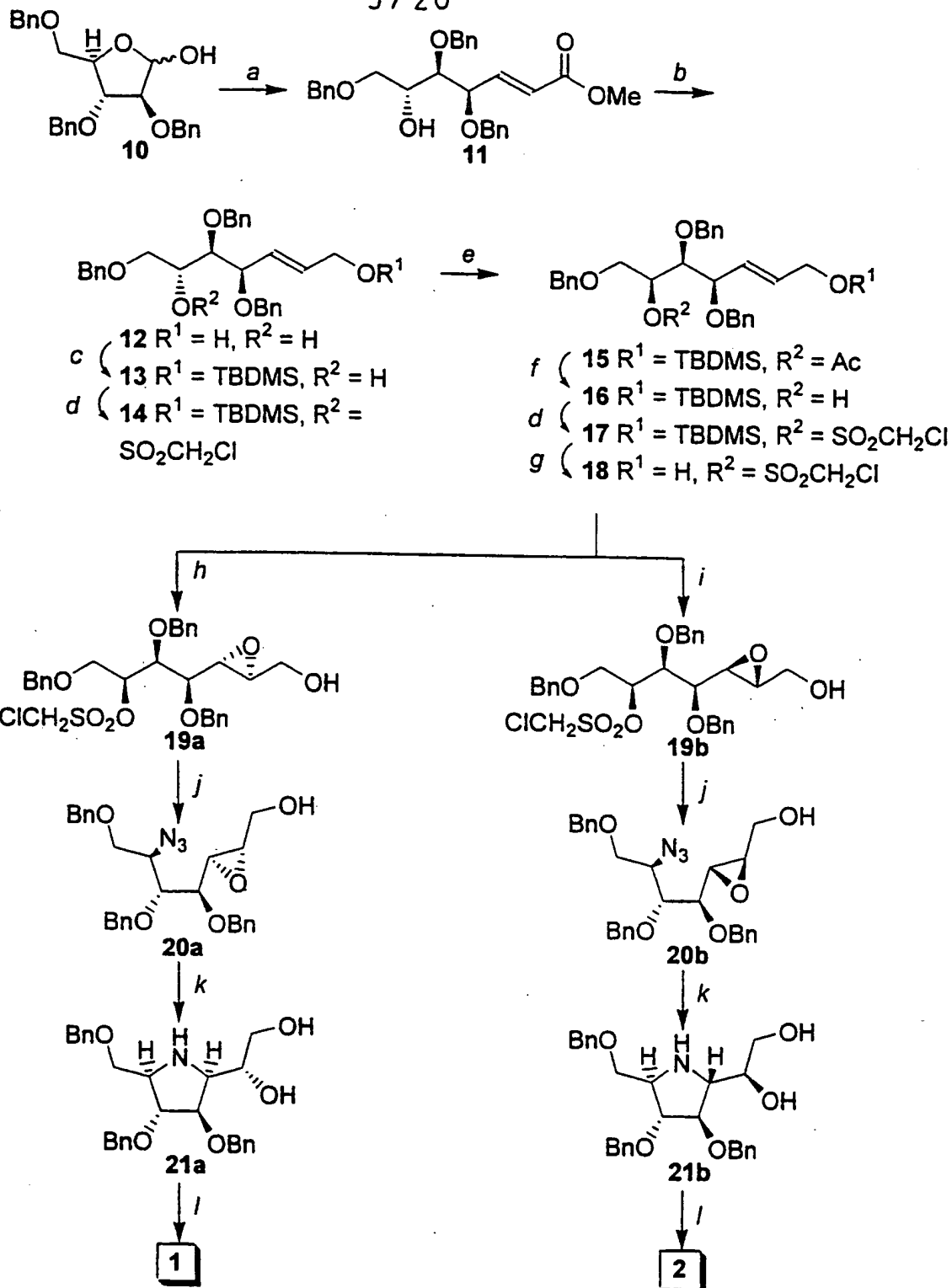


FIG. 5

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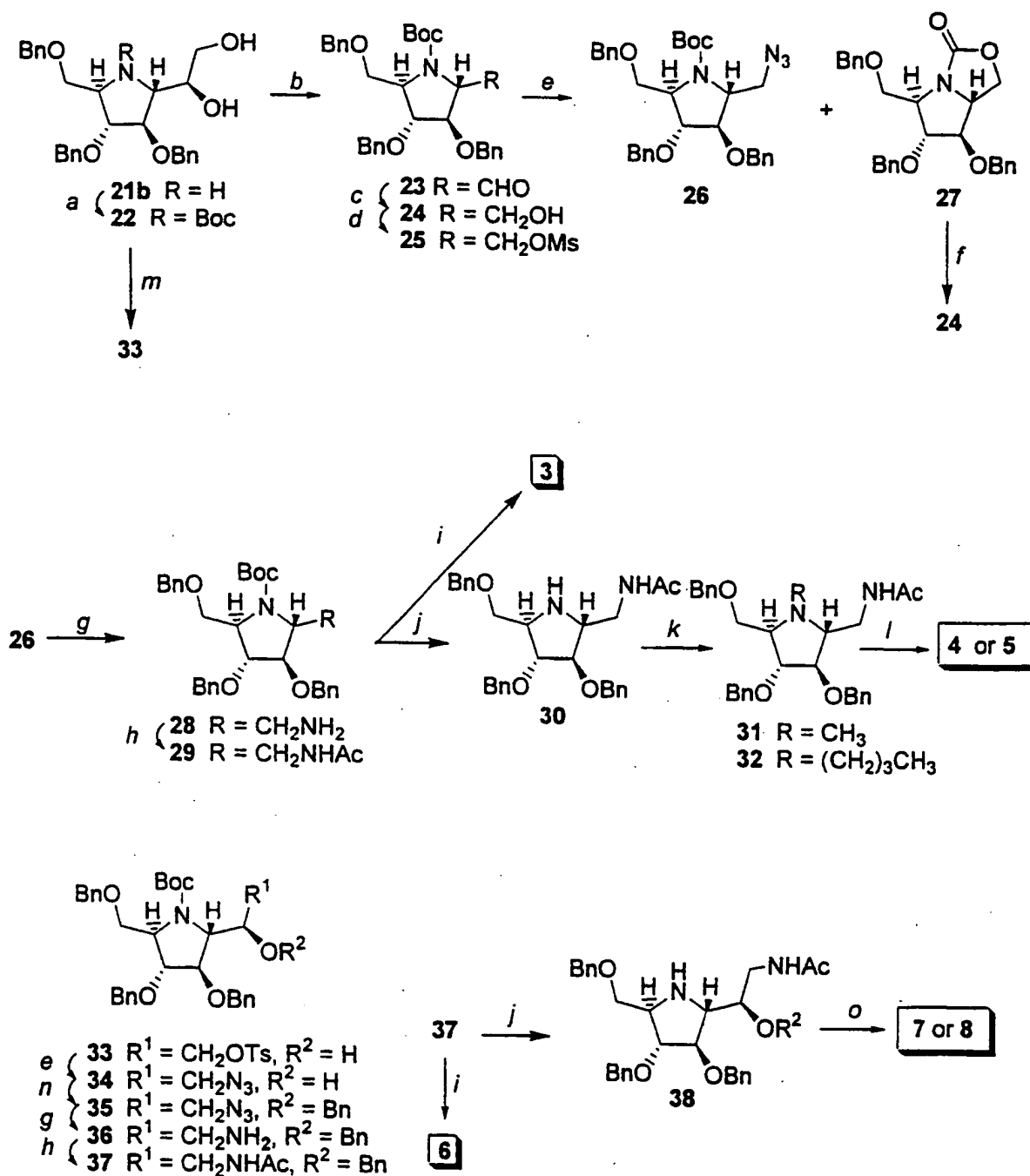
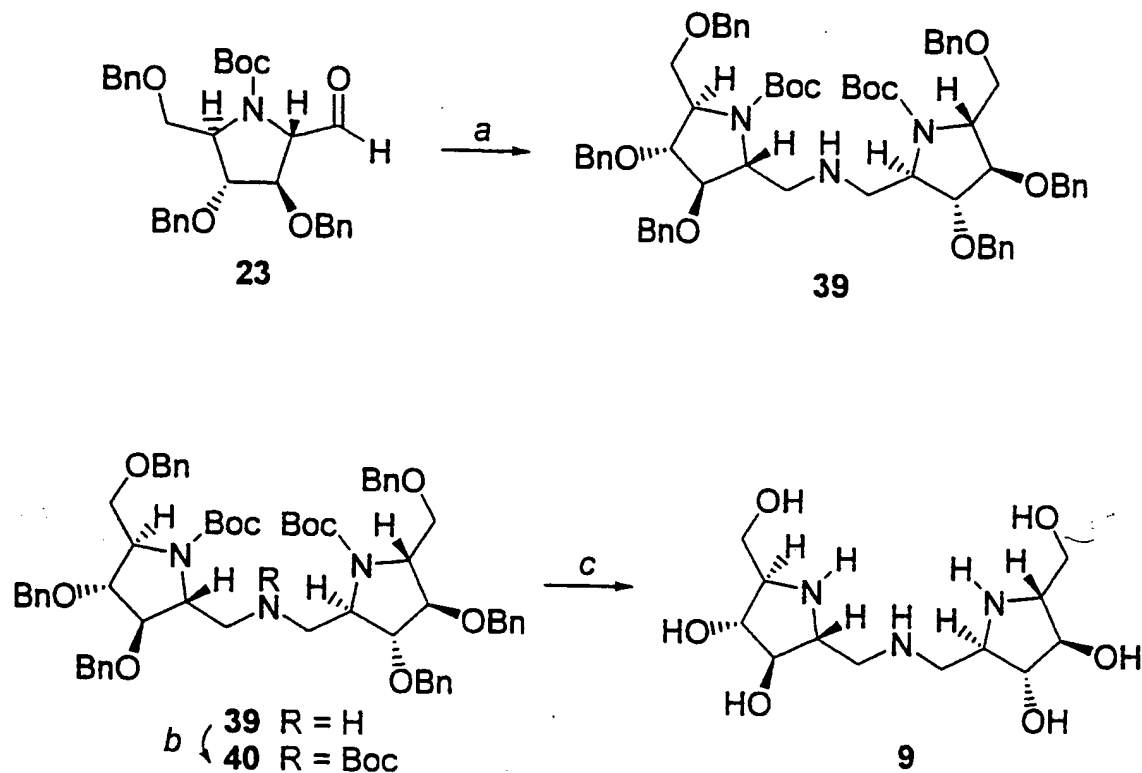


FIG. 6

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a $\text{NH}_4\text{OAc} - \text{NaBH}_3\text{CN} / \text{MeOH}$; *b* $(\text{Boc})_2\text{O} - \text{Et}_3\text{N} / \text{CH}_2\text{Cl}_2$; *c* 1) $\text{Pd/C} / \text{MeOH} - \text{HCl}$, 2) TFA.

FIG. 7

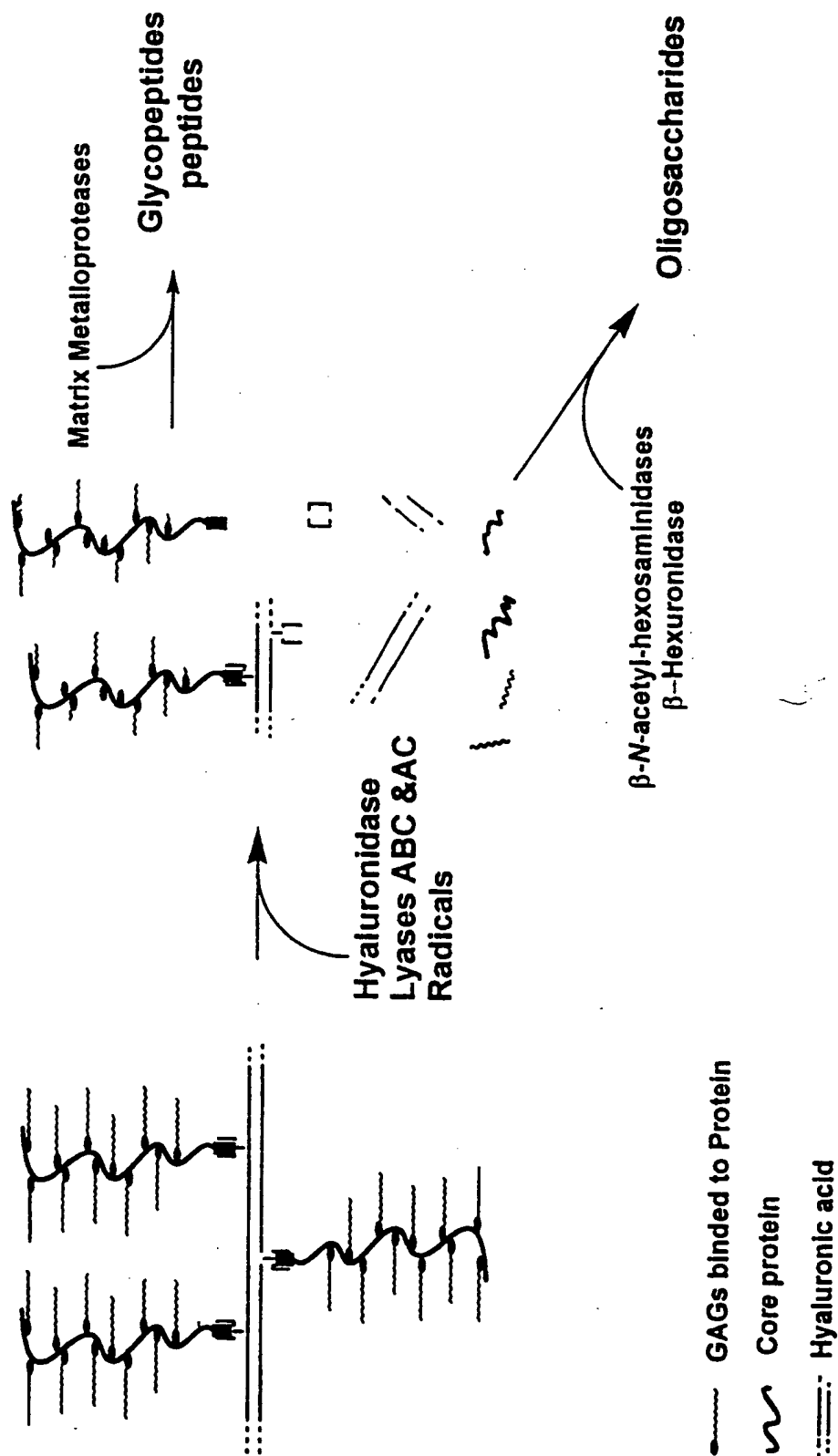


FIG. 8

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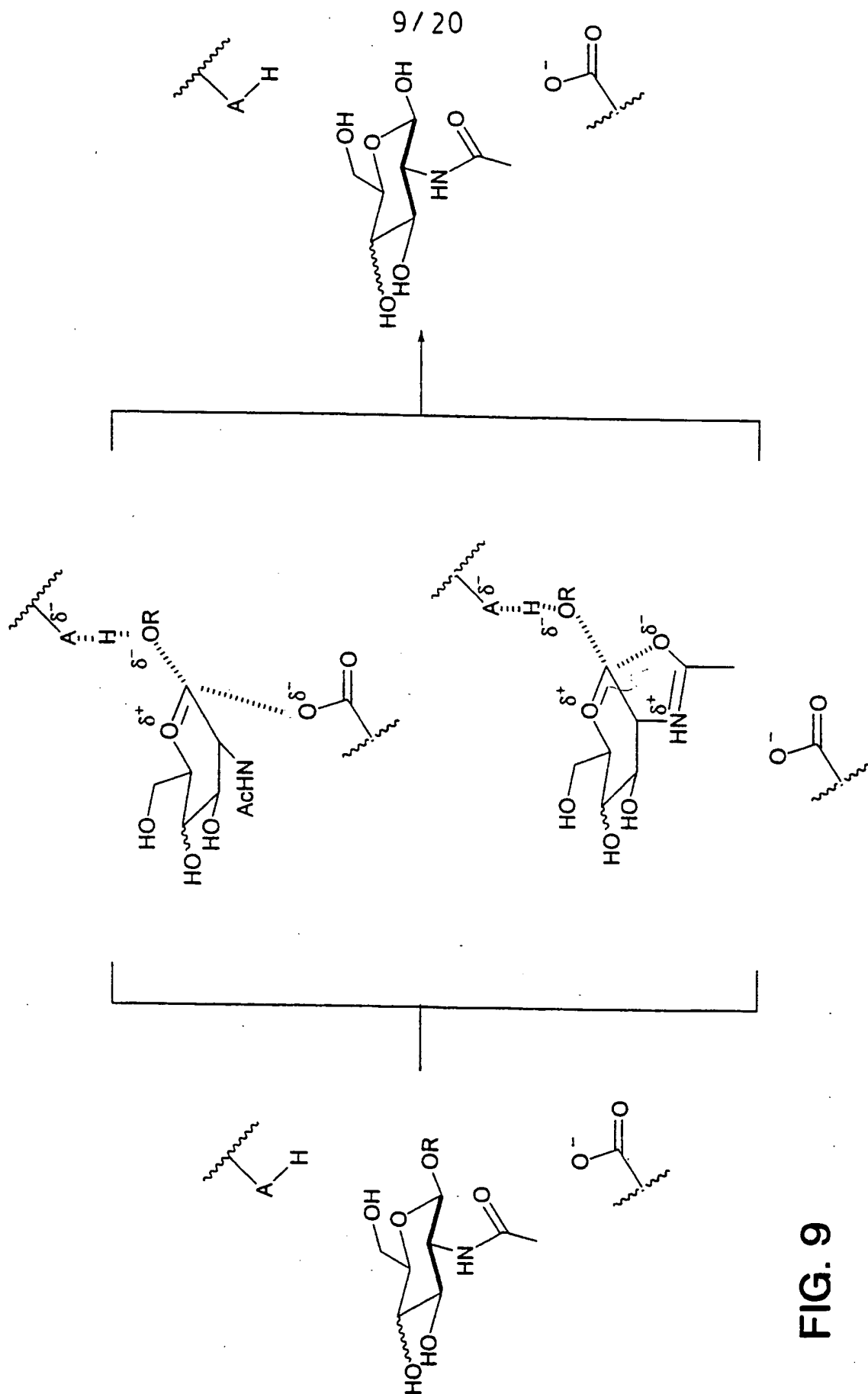


FIG. 9

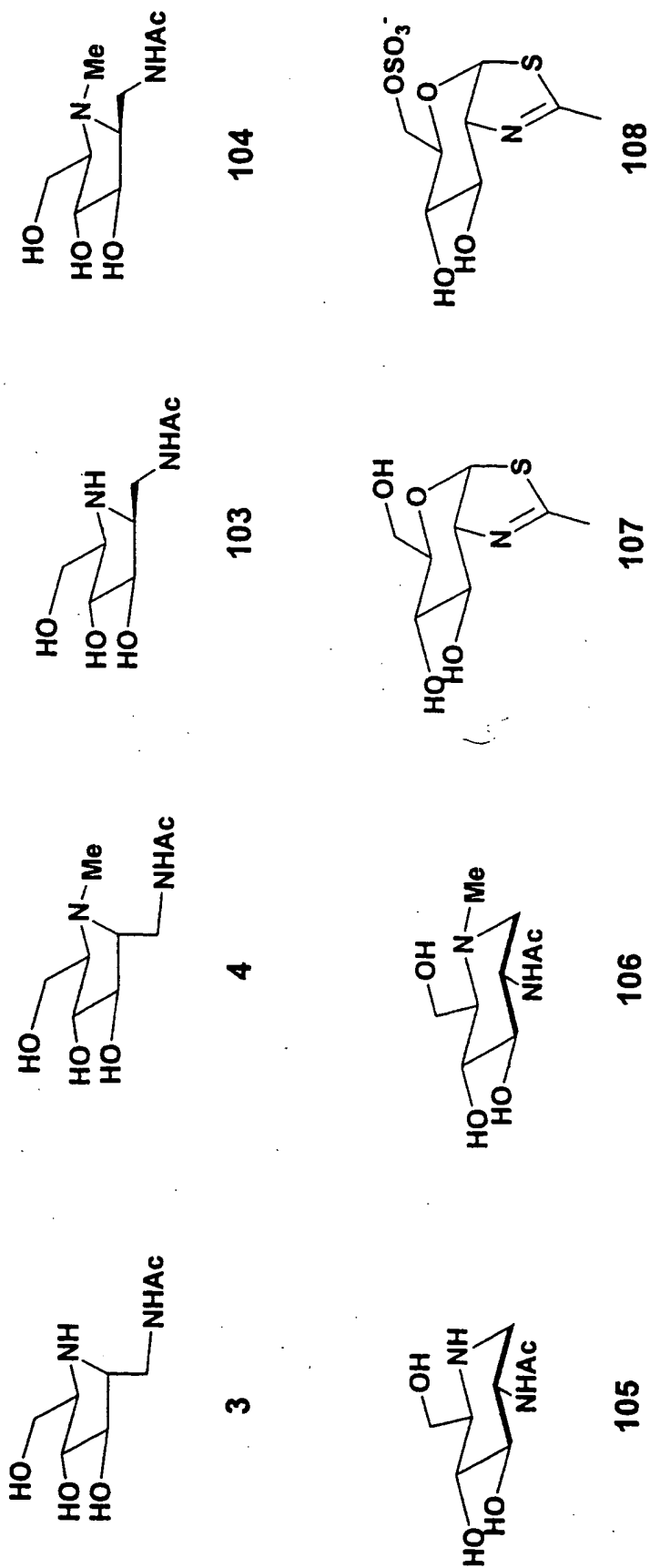
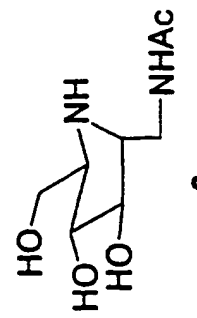
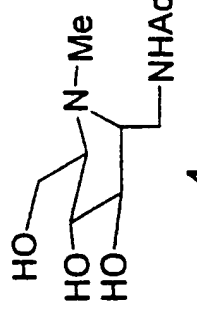
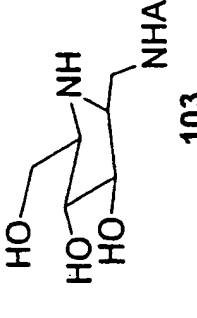
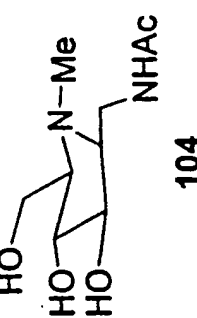
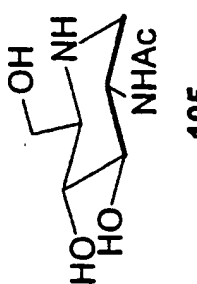
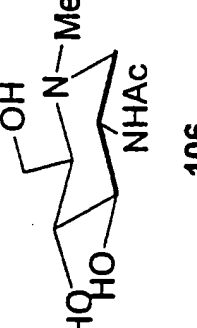
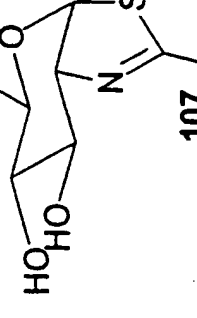
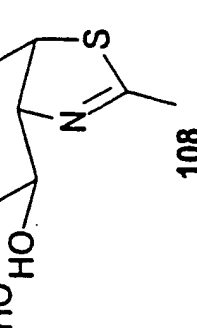


FIG. 10

 <p>3</p>	 <p>4</p>	 <p>103</p>	 <p>104</p>
Ki	—	24nM	—

 <p>105</p>	 <p>106</p>	 <p>107</p>	 <p>108</p>
Ki	1200nM	860nM	IC ₅₀ MUG=100μm IC ₅₀ MUGS<10μm

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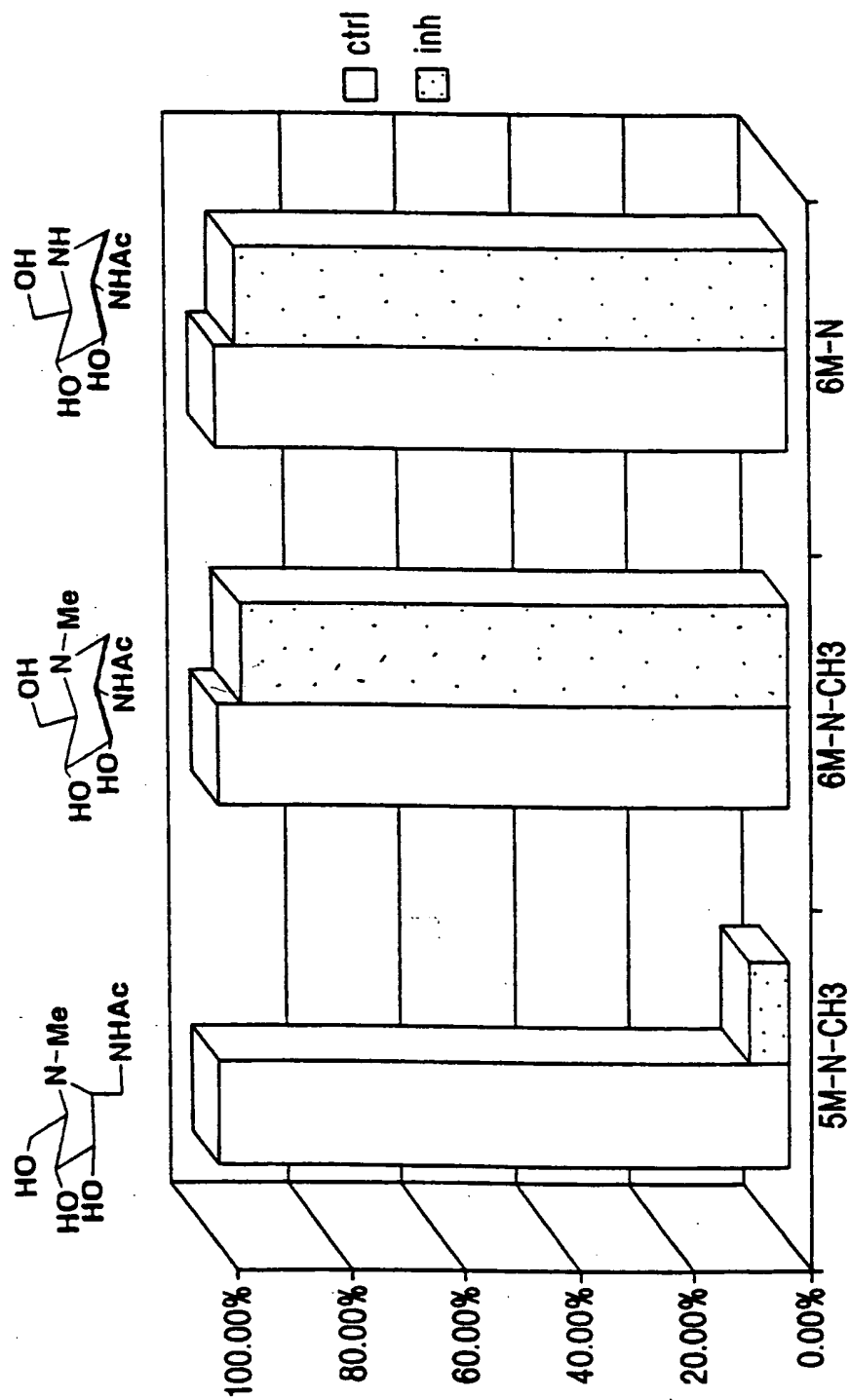
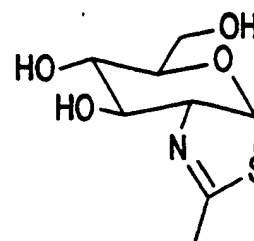
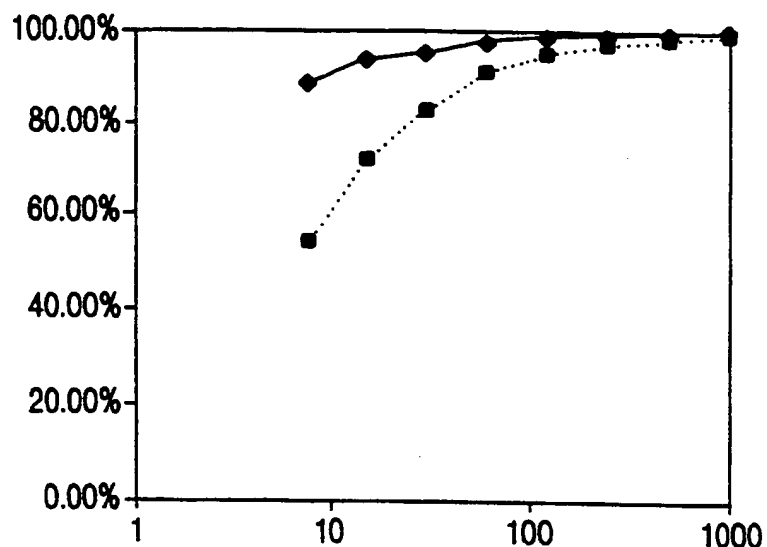
EFFECT OF SELECTED HEXOSAMINIDASE INHIBITORS ON INTRACELLULAR
HEXOSAMINIDASE ACTIVITY

FIG. 12

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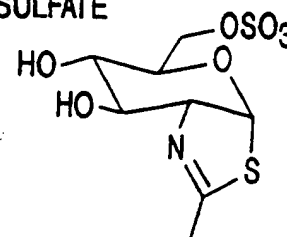
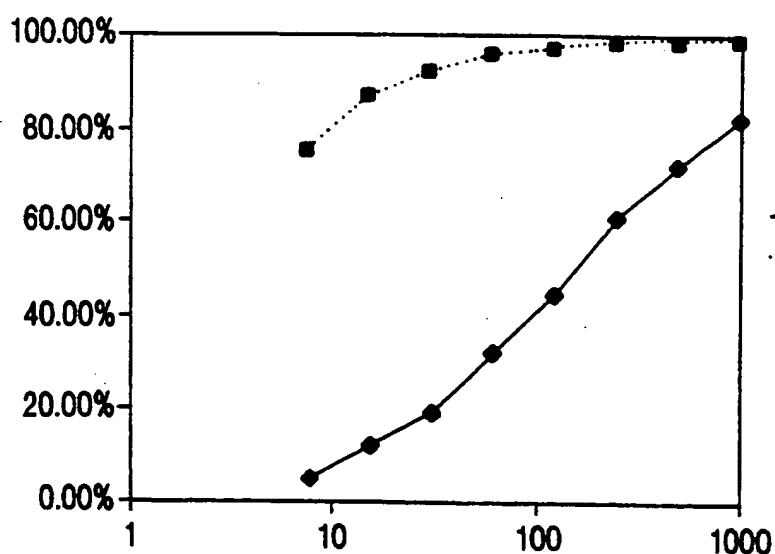
ENZYME - HUMAN PLACENTAL HEXOSAMINIDASE A
INHIBITOR - N - ACETYLGUCOSAMINE - THIAZOLINE



—●— MUG
- - -■- MUGS

FIG. 13A

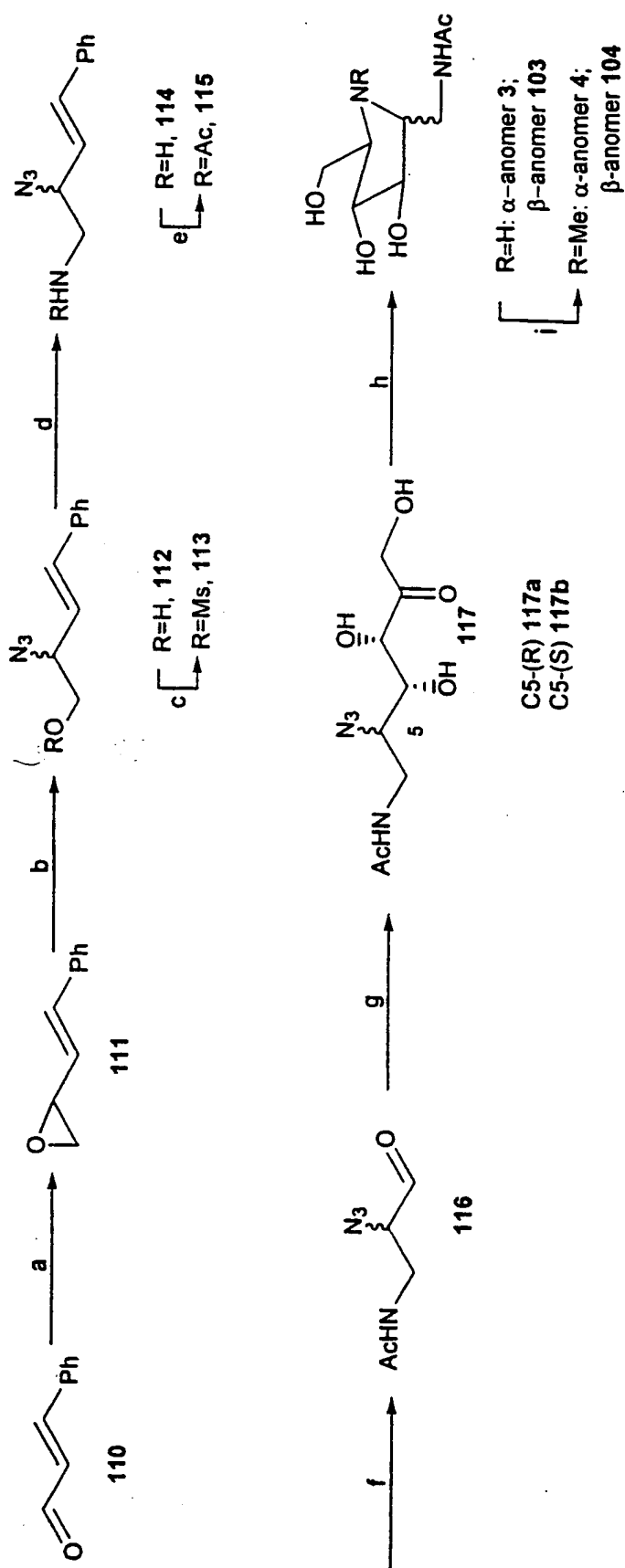
ENZYME - HUMAN PLACENTAL HEXOSAMINIDASE A
INHIBITOR - N - ACETYLGUCOSAMINE - THIAZOLINE - 6 SULFATE



—●— MUG
- - -■- MUGS

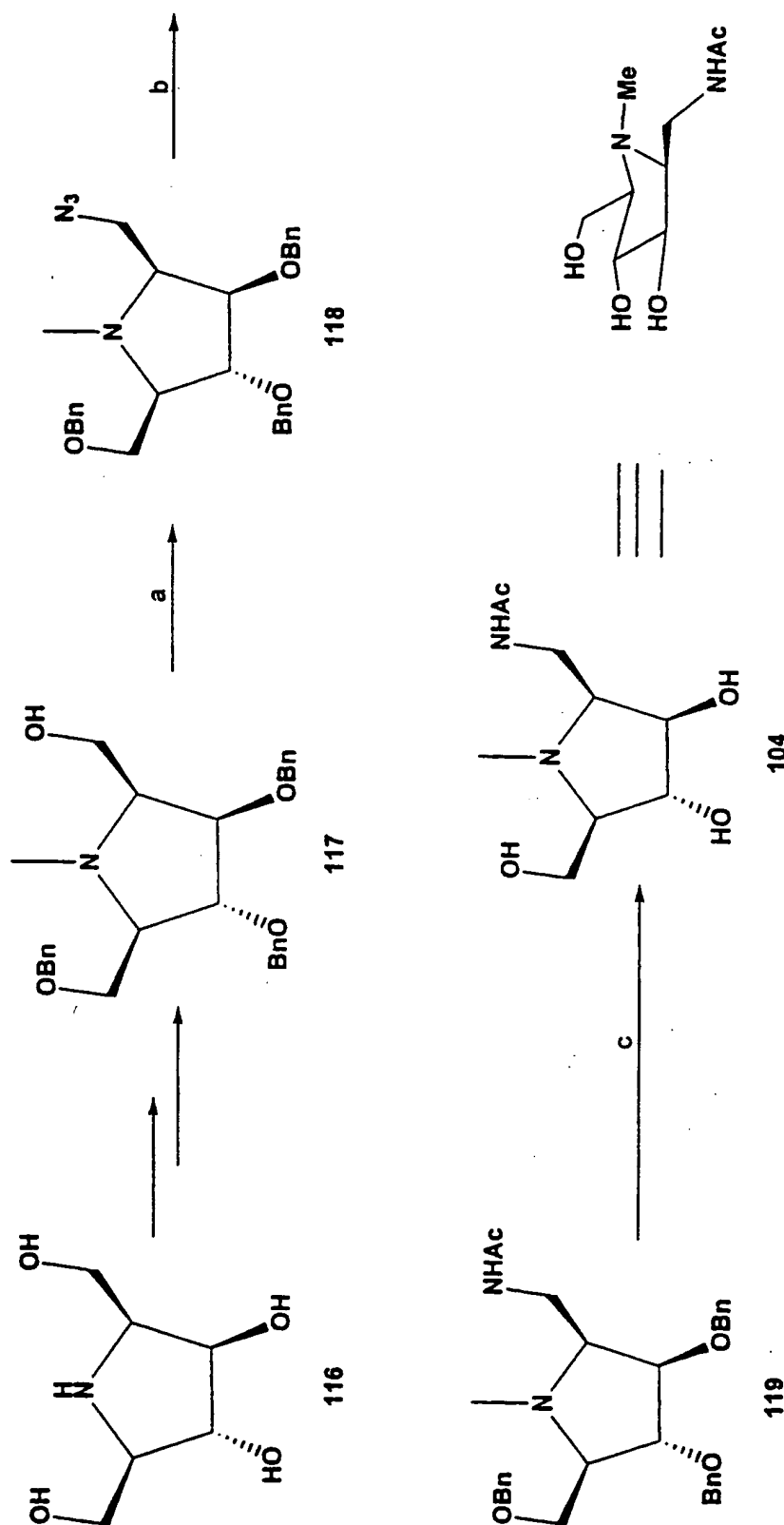
FIG. 13B

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a. $\text{Me}_3\text{S}^+\text{I}/\text{NaH}$, DMSO/THF; b. NaN_3 , acetone/ H_2O , 82% from 110; c. MsCl , Pyr. 96%; d. HMTA, NaI/EtOH ; HCl , 65°C ; e. isopropenyl acetate, 85% from 113; f. O_3 , Me_2S ; g. DHAP, RAMA, $\text{pH}=6.5$; acid phase 37°C , $\text{pH}=4.7$; 44% for (R), 30% for (S); h. $\text{Pd-C}/\text{H}_2$, 80%; i. CH_2O , $\text{Pd-C}/\text{H}_2$, 90%.

FIG. 14



a. MsCl, Pyr; NaN₃, CH₂Cl₂, 87% for 2 steps; b. PPh₃, THF; Ac₂O, Pyr, 87% from 118; c. Pd-C/H₂ 50 psi, 89%.

FIG. 15

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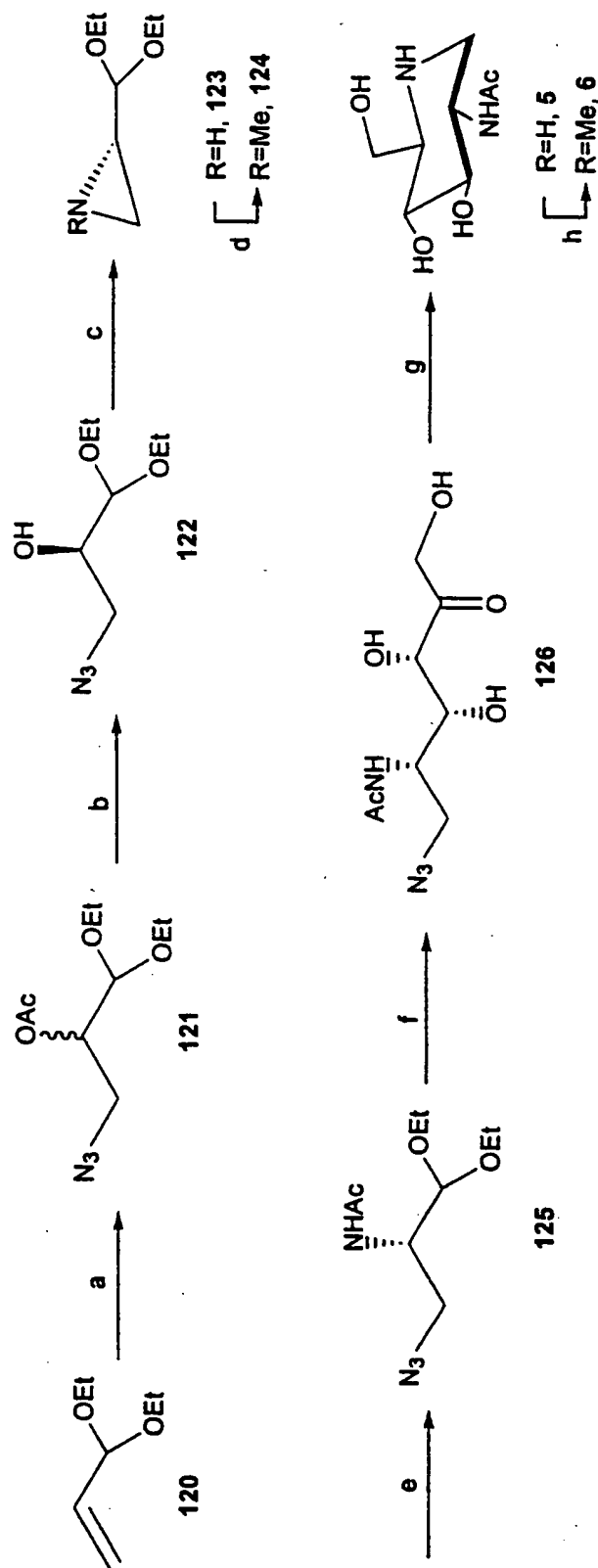
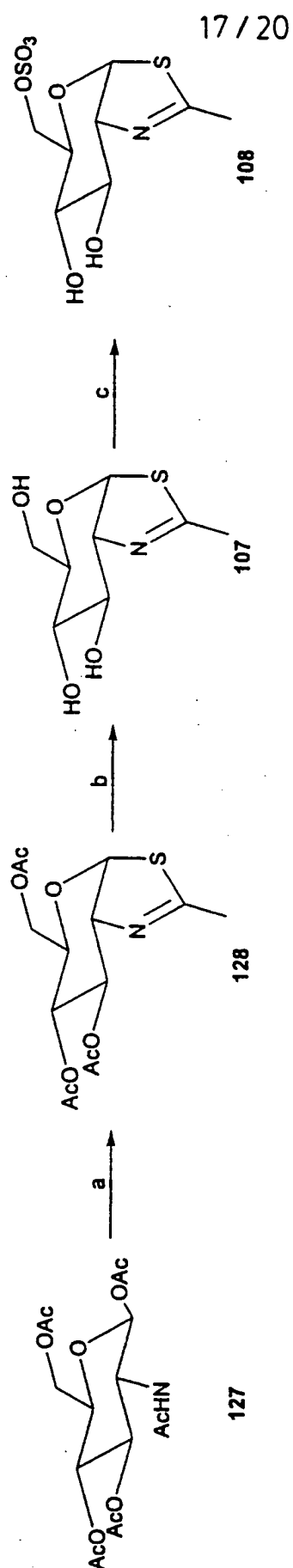


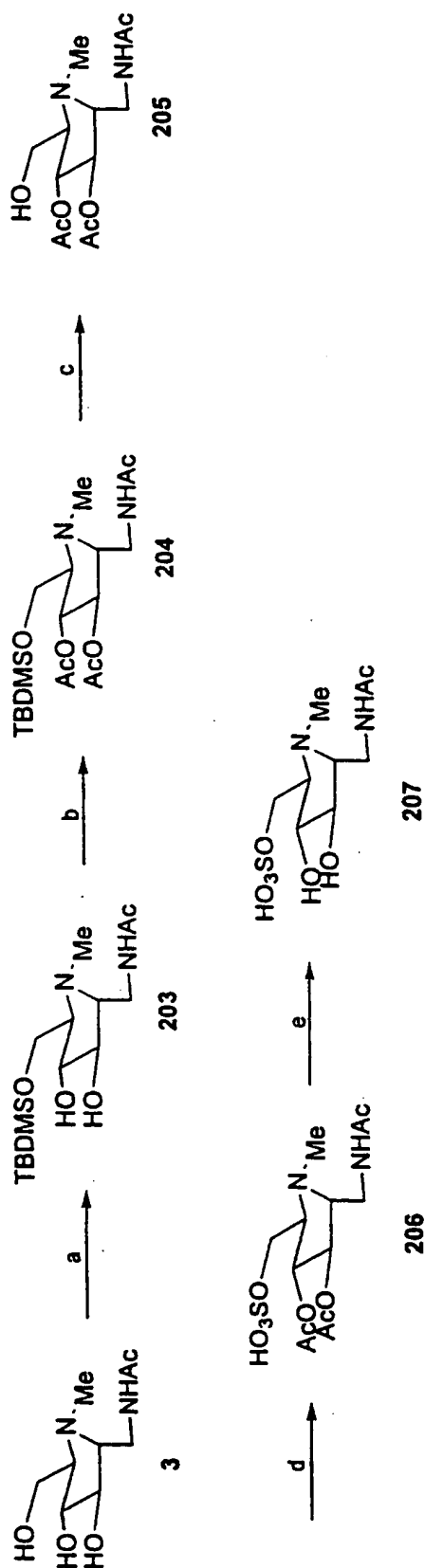
FIG. 16

a. H_2O_2 , PhCN , NaN_3 , $\text{pH}=7.5$; Ac_2O , Pyr , 76% for 3 steps; b. PS-80 , $\text{pH}=7.0$, 45%, 98% ee; c. Ph_3P , toluene, 120°C ; d. Ac_2O , K_2CO_3 , 30% for 2 steps; e. NaN_3 , $\text{ZnCl}_2/\text{Et}_2\text{O}$, DMF , 75°C , 62%; f. $\text{pH}=1$, 45°C ; DHAP , RAMA , $\text{pH}=6.5$; $\text{pH}=4.7$, acid phase, 37°C , 55% for 3 steps; g. $\text{Pd-C}/\text{H}_2$, 87%; CH_2O , $\text{Pd-C}/\text{H}_2$, 92%.



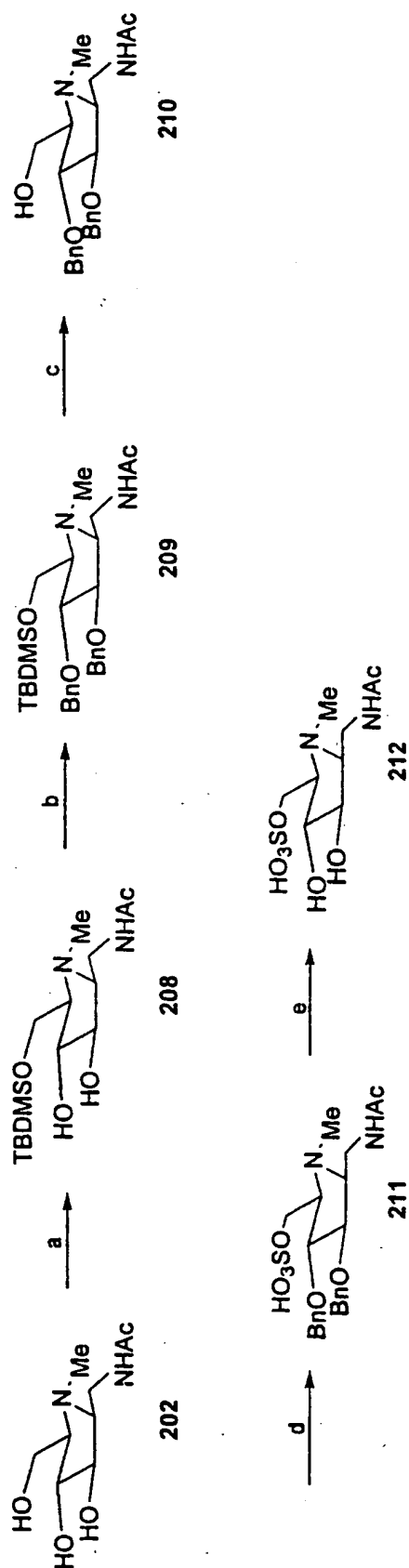
a. Lawesson's reagent, toluene, 80°C; b. MeONa/MeOH, 85% for 2 steps; c. SO₃·NMe₃, Pyr. 0°C, 87%.

FIG. 17



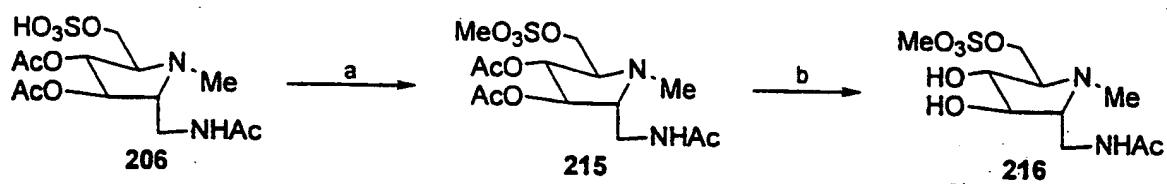
a. TBDMSCl, TEA, 0°C, DMF, overnight, 88%; b. Ac₂O, Pyridine, 0°C-rt.; c. AcOH/H₂O/THF(5:1:3), 50°C. overnight, 75% for two steps; d. SO₃/Pyr, pyridine, 25 °C. 82%; e. cat. MeONa, MeOH, 85%

FIG. 18



a. TBDMSOTf, TEA, 0 °C, DMF, 1.0 h, 90%; b. BnBr, NaH, 0 °C - 25 °C, 90%; c. TBAF, THF, 0 °C - 25 °C, 4h, 80%; d. SO₃/Pyr, pyridine, 25 °C, 80%; e. Pd(OH)₂/C, H₂, 75%

FIG. 19



a. MeOH, 50°C, 1h, 90%; b. MeONa (cat.), MeOH, 3h, 80%.

FIG. 20